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TITLE:

Preparation of trifluoromethyl steroids as postcoital

contraceptives

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Faming Zhuanli Shenqing Gongkai Shuomingshu, 17 pp.

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AB

[R H, He RI acetoxy, OH, CO2H, H; R2 = H, acetoxy, OH; R3 = H, OH; R4 = CF3, trifluorohydroxyalkyl; there may be double bonds in rings A or/and B] are prepd. Thus, 3.beta.-acetoxyandrost-5-en-17-one in THF contg. Me4NF was treated with CF3SiMe3 at room temp. for 3 h to give 83% 3.beta.-acetoxy-17.alpha.-(trifluoromethyl)androst-5-en-17.beta.ol. In a study using 6-days female rats, 17.alpha.-(trifluoromethyl)estra-1,3,5(10)-triene-3,17.beta.-diol (also prepd.) at 10 mg/Kg p.o. effected bleeding the day following the administration.

IT 161225-97-4P

> RL: BAC (Biological activity or effector, except adverse); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(prepn. of fluoromethyl steroids as postcoital contraceptives)

RN 161225-97-4 CAPLUS

CN Ergostane-3,6,24-triol, 28,28,28-trifluoro-, (3.alpha.,5.beta.,6.alpha.,24 .xi.) - (9CI) (CA INDEX NAME)

Absolute stereochemistry.

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